



4160-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2013-D-0575]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Guidance for Industry on Expedited Programs for Serious Conditions-Drugs and Biologics

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled "Guidance for Industry on Expedited Programs for Serious Conditions-Drugs and Biologics" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: FDA PRA Staff, Office of Operations, Food and Drug Administration, 1350 Piccard Dr., PI50-400B, Rockville, MD 20850, [PRStaff@fda.hhs.gov](mailto:PRStaff@fda.hhs.gov).

SUPPLEMENTARY INFORMATION: On December 31, 2013, the Agency submitted a proposed collection of information entitled "Guidance for Industry on Expedited Programs for Serious Conditions-Drugs and Biologics" to OMB for review and clearance under 44 U.S.C. 3507. An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910-0765. The approval expires on March 31, 2017. A copy of the supporting statement for this

information collection is available on the Internet at

<http://www.reginfo.gov/public/do/PRAMain>.

Dated: April 25, 2014.

Leslie Kux,

Assistant Commissioner for Policy.

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